P10A/REV05

TRANSMITTAL OF INFORMATION DISCLOSURE STAFFMENT (Under 37 CFR 1.97(b) or 1.97(c))						Docket No. 17719 (PC27263A)	
In Re Application: Manuela Villa, et al.							
Application No.	Filing Date	Examiner	FT 8	Eustomer No.	Group Art Unit	Confirmation No.	
10/522,254	January 25, 2005	Unassigne	d	23389	Unassigned	Unassigned	
Title: BICYCLO-PYRAZOLES ACTIVE AS KINASE INHIBITORS, PROCESS FOR THEIR PREPARATION AND PHARMACEUTICAL COMPOSITIONS COMPRISING THEM							
	(Only co	Payme mplete if Applicant elect	ent of Fee s to pay the f	fee set forth in 37	CFR 1.17(p))		
 □ A check in the amount of is attached. ☑ The Director is hereby authorized to charge and credit Deposit Account No. 19-1013/SSMP as described below. □ Charge the amount of ☑ Credit any overpayment. ☑ Charge any additional fee required. □ Payment by credit card. Form PTO-2038 is attached. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038. 							
l certify that this account is being and Trademark (Date)	Signature Printed Name of Person Sig	ntion to charge deposition to	I hereby or the United class mai Patents, P 1.8(a)] on Ma	ertify that this correction that this correction in an envelope of the correction of	spondence is being vice with sufficient paddressed to "Corrandria, VA 22313-1 rson Mailing Correspondence in the corporation of Person Mailing Corporation in the corporation of Person Mailing Corporation in the corporation	deposited with ostage as first unissioner for 450" [37 CFR	
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Manuela Villa, et al.

Examiner:

Unassigned

Serial No.:

10/522,254

Art Unit:

Unassigned

Filed:

January 25, 2005

Docket:

17719 (PC27263A)

For:

BICYCLO-PYRAZOLES ACTIVE AS

Dated:

March 3, 2005

KINASE INHIBITORS, PROCESS FOR THEIR PREPARATION AND PHARMACEUTICAL

COMPOSITIONS COMPRISING THEM

Confirmation No.: Unassigned

Mail Stop Amendment Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

INFORMATION DISCLOSURE STATEMENT

Sir:

In accordance with 37 C.F.R §§1.97 and 1.98, it is requested that the following references, which are also listed on the attached Form PTO-1449, be made of record in the above-identified case.

- 1. United States Patent Application Publication No. 2003/0073672 A1, published April 17, 2003 to Breitenbucher et al.;
- 2. PCT International Publication No. WO 02/064574 A2, published August 22, 2002;
- 3. PCT International Publication No. WO 02/12242 A2, published February 14, 2002;

CERTIFICATE OF MAILING UNDER 37 C.F.R. §1.8(a)

I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Mail Stop Amendment, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Dated: March 3, 2005

Peter I Bernstein

- 4. PCT International Publication No. WO 96/12720, published May 2, 1996;
- 5. Kikuchi C. et al., "Tetrahydrothienopyridylbutyl-Tetrahydrobenzindoles: New Selective Ligands of the 5-HT₇ Receptor", *Bioorganic & Medicinal Chemistry Letters*, 12(18):2549-2552 (2002), XP-002256018;
- 6. Singh P. et al., "Quantitative Structure-Activity Relationship Studies on a New Class of Antihypertensive Agents: Derivatives of 3-Aryl-4,5,6,7-Tetrahydro-1H-Pyrazolo[4,3-c]Pyridine", Quantitative Structure-Activity Relationships, 9(1):29-32 (1990), XP-001155089;
- 7. Winters G. et al., "Synthesis, in Vitro [³H]Prazosin Displacement, and in Vivo Activity of 3-Aryl-4,5,6,7-Tetrahydropyrazolo[4,3-c]Pyridines, a New Class of Antihypertensive Agents", *J. Med. Chem.*, 28(7):934-940 (1985), XP-002256019;
- 8. Radinov R. et al., "3-Phenylpyrazolo[4,3-c]Pyridine and Derivatives: Structure Determination", *Journal of Molecular Structure*, 158:99-108 (1987), XP-009018257;
- 9. Lackey K. et al., "The Discovery of Potent cRaf1 Kinase Inhibitors", Bioorganic & Medicinal Chemistry Letters, 10(3):223-226 (2000), XP-004188821; and
- 10. Philip Cohen, "The Development and Therapeutic Potential of Protein Kinase Inhibitors", *Current Opinion in Chemical* Biology, 3(4):459-465 (1999), XP-002216616.

The references were cited in a Search Report dated October 27, 2003 received from the European Patent Office. Applicants are submitting copies of the above-cited references, together with a copy of the Search Report. The relevance of the above-identified references has been described in the Search Report.

Inasmuch as this Information Disclosure Statement is being submitted in accordance with the schedule set out in 37 C.F.R. §1.97(b), no statement or fee is required.

Respectfully submitted,

Peter I. Bernstein

Registration No. 43,497

Scully, Scott, Murphy & Presser 400 Garden City Plaza-Suite 300 Garden City, New York 11530 (516) 742-4343

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•				Manuela Villa, et al.		·		
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		WO 02/064574 A2	8/22/02	PCT				
	'	WO 02/12242 A2	2/14/02	PCT				
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			OTHER	DOCUMENTS (Including As	ithor, Title, I	Date, Pertinent l	ages, Etc.)	
A b.	Kikuchi C. et al., "Tetrahydrothienopyridylbutyl-Tetrahydrobenzindoles: New Selective Ligands of the 5-HT ₇ Receptor", <i>Bioorganic & Medicinal Chemistry Letters</i> , 12(18):2549-2552 (2002), XP-002256018							
	Singh P. et al., "Quantitative Structure-Activity Relationship Studies on a New Class of Antihypertensive Agents: Derivatives of 3-Aryl-4,5,6,7-Tetrahydro-1H-Pyrazolo[4,3-c]Pyridine", Quantitative Structure-Activity Relationships, 9(1):29-32 (1990), XP-001155089							
	Winters G. et al., "Synthesis, in Vitro [3H]Prazosin Displacement, and in Vivo Activity of 3-Aryl-4,5,6,7-Tetrahydropyrazolo[4,3-c]Pyridines, a New Class of Antihypertensive Agents", J. Med. Chem., 28(7):934-940 (1985), XP-002256019							
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	LIST OF PRIOR ART CATALRANGE BY APPLICANT	17719 (PC27263A)	10/522,254			
(Us	se several sheets if necessary)					
• •		Applicant(s) Manuela Villa, et al.				
•		Filing Date January 25, 2005	Group Art Unit Unassigned			
	OTHER	DOCUMENTS (Including Author, Title, I	<u> </u>			
	Radinov R. et al., "3-Phenylpyrazol of Molecular Structure, 158:99-108	o[4,3-c]Pyridine and Derivatives: Struc (1987), XP-009018257	cture Determination", Journal			
	Lackey K. et al., "The Discovery of Potent cRaf1 Kinase Inhibitors", Bioorganic & Medicinal Chemistry Letters, 10(3):223-226 (2000), XP-004188821					
	Philip Cohen, "The Development and Therapeutic Potential of Protein Kinase Inhibitors"; Current Opinion in Chemical Biology, 3(4):459-465 (1999), XP-002216616					
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